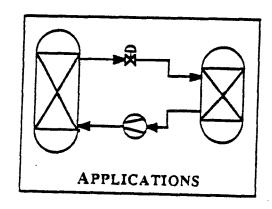
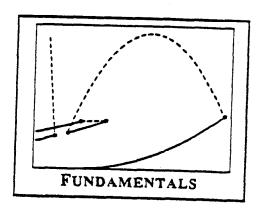
PROCEEDINGS

2ND INTERNATIONAL SYMPOSIUM ON SUPERCRITICAL FLUIDS





SUPERCRITICAL CARBON DIOXIDE EXTRACTION OF SPENT BLEACHING CLAYS

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ABSTRACT

The disposal of spent bleaching clay generated in the vegetable oil processing industry is a potential problem of environmental concern, due the pyrophoric nature of the residual oil/clay mixture in landfills. This poster describes the use of SC-CO2 for processing both neutral and acidic clays used in the refining of soybean oil. Rapid extractions of spent bleaching clay can be affected at 10,000-12,000 psig by proper preparation of the clay substrate. Experiments utilizing both laboratory and pilot plant extractors have yielded almost 100% of the adsorbed oil. Extraction of the clay can be facilitated by crossblending the clay with a distomaceous earth-based dispersant. The oils recovered by the supercritical fluid extraction (SFE) process have properties (free fatty acid content, Lovibond color ratio, and phosphorus content) similar to those found for degummed-bleached oils.

INTRODUCTION

An average soybean oil refinery generates about 5,000 lbs. of spent bleaching clay per day in the removal of odoriferous components and color bodies from refined feedstock. This processing aid must be treated to remove the adsorbed residual oil or be buried in a landfill after the bleaching step (1). Several processes have been developed to strip residual oil from the clay matrix based on treatment of the clay with dilute caustic solution (2) or extraction of the oil using organic solvents (3). These processes also produce unacceptable chemical residues which can impact negatively on the environment (4).

Currently, spent bleaching clays from the vegetable oil processing industry are disposed of in landfills (5), necessitating additional costs for transport of the clay from the refinery and its subsequent treatment at the fill site. At the landfill, the clay is frequently buried or treated with water, to reduce the probability of spontaneous combustion of the clay/oil mixture. With the exception of the recent study by Taylor and Jenkins (6), little is known about the factors governing the pyrophoricity of spent bleaching clays.

Treatment of spent bleaching clays with supercritical carbon dioxide (SC-CO2) presents an alternative to the above disposal methods. SFE utilizing carbon dioxide provides an environmentally acceptable solution for post refinery treatment of the clay, since CO2 is nonflammable, nontoxic, and is a selective solvent for oleophilic materials. One study has been reported (7) in which supercritical pro-

pane and fluorocarbons were utilized in the treatment of spent clays imbibed with silicone and vegetable oils. Unfortunately, in this work, only one experiment was performed on a clay/vegetable oil mixture, resulting in a low recovery of the oil. For this reason, and to avoid the use of potentially explosive and environmentally unacceptable supercritical fluids, we initiated an extraction study using SC-CO2.

EXPERIMENTAL

Successful extraction of spent bleaching clays with SC-CO2 can be achieved only by optimizing the experimental conditions. Rapid extraction of the oil from the clay was facilitated by conducting the extractions at 12,000 psig and 80°C. a condition where the oil solubility in SC-CO2 is maximized (8,9). The particle size of bleaching clays can also inhibit complete extraction of the oil from the matrix, due to channeling of extraction fluid through the sorbent bed (7), as well as compaction of the clay bed under high pressure. For this reason, studies were conducted on both a laboratory, and a pilot plant scale. The pilot plant study utilized a captive CO2 supply and was operated in a recyle mode, thereby permitting simulation of an industrial scale process.

Typical conditions utilized in the clay extraction studies are tabulated in Table 1. The laboratory-scale extraction unit has been previously described (9). Bleaching clay was poured into the specified tubular extraction vessel and extraction commenced at 11,000 psig and 80°C. Extractions were continued until oil could no longer be removed from the receiver attached to the extraction system.

Table 1

TYPICAL CONDITIONS FOR LAB AND PROT PLANT EXTRACTIONS

Pameter	Laboratory	Plot Plant	
Extraction Voscol	Tubular (2' x 6/8" LDJ	4-Liter Vessel (2° x 2° LDJ)	
Extraction P & T	11,000 paig, 80G	12,000 peig, 80C	
Collection P & T	Ambient	2,300 paig, 80C	
CO2 Used	1.5 - 4.1 Be.	6 - 8 Mg.	
Clay Weight	0.09 - 0.25 ba.	6.1 - 8.6 Mm	

Pilot plant extractions were performed on the NCAUR semi-continous supercritical fluid extraction unit shown in Figure 1. Details on the construction of the extractor, which has provision for recycling the extraction fluid, can be found in the literature (10). Clay extractions were performed in one of the 4-liter extraction vessels (A) shown in the schematic diagram. Extractions were conducted in this vessel at 12,000 psig and 80°C using a CO2 flow rate of approximately 0.5 lbs./min. The dissolved soybean oil in the SC-CO2 was

transported to the receiver vessel (R), which was operated at 2,300 psig and 40°C. Under these conditions, soybean oil exhibits less than 0.1 wt % solubility in the compressed CO2, and readily precipitates from the fluid phase. Extractions could be completed under the above conditions on 5.0-8.5 lbs. charges of clay using 6-8 lbs. of CO2 that was then recycled over the clay.

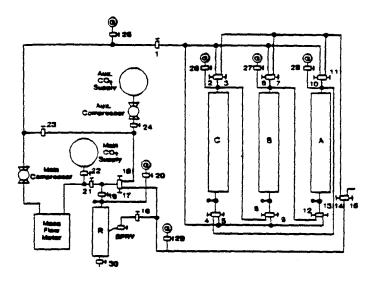


Fig. 1 - Semi-continuous supercritical fluid extraction system.

The spent bleaching clays utilized in these studies were obtained from refineries operated by Riceland Foods (Stuttgart, AK) and Central Soya (Ft. Wayne, IN). Both neutral and acidic clays were incorporated in this study to test the effectiveness of SC-CO2 extraction with respect to the clay structure. A pelletized distomaceous earth, designated Hydromatrix, from Analytichem International (Harbor City, CA) was used in several experiments as an aid in dispersing the spent bleaching clay.

Samples of both the extracted clay and recovered oil were saved from each experiment. Selected oil samples were characterized with respect to their free fatty acid content (FFA), peroxide value (PV), Lovibond color ratios, and phosphorus content. Standard AOCS Methods (11) were used in performing these tests unless otherwise noted. The Lovibond color ratios of the extracted oils were determined using a computerized Colourscan system while the phosphorus content of the oils were analyzed by atomic absorption spectroscopy. Oil content of the clay was assessed by Soxhlet extraction with n-hexane.

RESULTS AND DISCUSSION

Experimental extraction results performed on both neutral and acidic bleaching clays are summarized in Table 2. Initial experiments were conducted on the neutral bleaching clay matrix using the pilot plant extractor. Extractions were terminated when oil could no longer be collected in the receiver vessel of the pilot plant. The oil recovered in Run 1, which is

expressed as a weight percent of the total clay charge, was 15.2%. The extracted clay in this experiment was removed from the extraction vessel and analyzed for oil content. The residual oil level was found to be 8.8 wt.% based on the initial clay charge. The results from this run indicated that either the extraction conditions or clay preparation were inadequate to effect total removal of the oil from the clay matrix.

SFE EXPERIMENTS ON SPENT BLEACHING CLAYS

Table 2

ALIN	Clay Type	Extractor	% Of Extraoted	Clay Pres
1	Neutral	Pflet	15.2%	
2	Neutral	Pliet	19.3%	***
3	Neutral	Pliot	26.8%	Mixed
4	Neutral	Pflot	25.5%	Mixed
4	Neutral	هما	32.1%	
•	Neutral	Leb	30.7%	Hydrometriz
7	Newtral	Pflot	30.9%	Hydrometrix
8	Acid	Leb	34.1%	
•	Apid	Plot	22.6%	
10	Nautrai	Pliet	26.6%	

. Total WL % of OE in Clay on 32-34%

An identical extraction (Run 2) was then performed to verify the results obtained in the first experiment. The oil yield in this case was found to be 19.3 wt.X. Oil analysis on the extracted clay yielded a 7.2 wt.X residual oil content, verifying that the extraction was not completed. These two initial results suggested that perhaps the oil distribution in the spent bleaching clay was not homogeneous and that mixing the substrate prior to extraction would enhance the efficiency of the extraction.

Two extractions (Runs 3 & 4) were then performed on the pilot plant extractor to verify the above hypothesis. Oil yields for these two experiments were 26.8 and 25.5 wt.%, respectively. Analysis of the SC-CO2 extracted clays for residual oil content showed that considerable present in the oil VAS still extracted clays (Run 3 - 6.12 wt.X, Run 4 - 8.8 wt.%). A discrepancy was also apparent in the total oil mess balance, since combining the extracted and residual oil wt.% for Runs 1, 2, and 4 yielded 24.0, 26.5, 32.9, and 34.3 wt.X, respectively. It was obvious that premixing the spent bleaching clay did not assure total extraction of all available oil.

Determination of the total oil content in spent clay matrix via Soxhlet extraction is at best difficult due to the fine particle size of clay and its tendency to agglomerate in the Soxblet extraction device. Accurate determination of the total oil content of the nest spent bleaching clay is essential in order to assess the extraction efficiency of the SPE process. Several exhaustive extractions of the starting clay material were made with the aid of a dispersing agent to establish the residual oil content of the sorbent. In this procedure, the spent bleaching clay was mixed in a weight ratio of 2:1 with pelletized distoraceous earth (Hydromatrix) to disperse the clay sample. This modification of the Soxhlet procedure facilitated contact between the clay matrix and extraction solvent and yielded consistent analytical results. The Soxhlet extraction values on the samples used in Runs 2, 3, and 4 were determined to be 31.7, 31.1, and 32.8 wt.X. The accuracy of these figures was further verified by running an extraction of spent bleaching clay in the small laboratory tubular extractor (Run 5) which yielded 32.1 wt.X of oil based on the initial clay charge. Soxhlet extraction of this clay sample gave 32.0 wt.X oil, a value in excellent agreement with the result obtained from SC-CO2 extraction and the previously quoted Soxhlet-derived data. These results suggest that SC-CO2 extraction could be used as an alternative to Soxhlet extraction with liquid hydrocarbon solvents.

The consistent results obtained by using the Hydromatrix dispersant in the modified-Soxhlet procedure suggested that the use of a dispersion agent in the SFE of clays would enhance the recovery of the soybean oil. Consequently an experiment was run using the laboratory scale extractor loaded with 40 grams of the spent clay plus 20 grams of Hydromatrix. The total yield of oil from this experiment (Run 6) was 30.7%. In addition, it was noted that the inclusion of Hydromatrix decreased the time of the extraction by 1/3 and required less than half the extraction fluid needed for the SFE of the nest clay. This trend is shown in Figure 2. Compaction of the spent bleaching clay bed in the extractor, with the attendant large pressure drop, is also avoided by using the dispersant for the SFE.

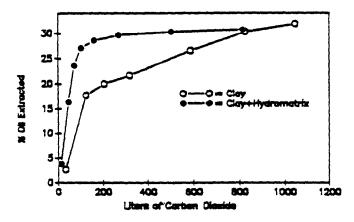


Fig. 2 - Comparison of oil extraction rates for clay vs. clay + Hydromatrix.

A similar experiment to the previous one was performed in the 4-liter pilot plant vessel using a 2:1, clay to Hydrometrix charge. In this case, a 30.7 wt.% oil yield was obtained after recycling the CO2 through the extractor bed. This result is in excellent agreement with the value obtained on the small laboratory extractor and exceeds the yields recorded on nest clay beds (Runs 1-4). This result was further verified by running another SFE on the nest clay in a 4-liter extraction vessel (Run 10). The yield of only 26.5 wt.% oil, clearly confirms the enhancement of oil yield by incorporation of the dispersant into the SFE scheme. Additional pilot plant runs were performed on an acidic spent bleaching clay from another industrial source. Extraction of this clay in the laboratory unit (Run 8) gave a 34.1 wt.%

yield. This result compared favorable with the Soxhlet-measured oil content of 34.6 wt.X. A pilot plant run was also made on the acidic clay, and produced a oil yield of 32.6 wt.X (Run 9). It therefore appears, that the SC-CO2 extraction of spent bleaching clays is equally applicable to both neutral and acidic matrices.

Several physical and chemical tests were run on selected oil samples taken from various experimental runs. The results of these tests, which included measurement of the oil's free fatty acid content, peroxide value, Lovibond color ratio, and phosphorus content, are listed in Table 3. Overall, the results in Table 3 indicate that the oils recovered from the SFE of spent bleaching clays, are approximately equivalent to a conventional degummed-bleached oil (12). Hence, these oils would be suitable for a number of industrial applications that do not require food grade oil specifications. Further refining of the SFE-derived oils would be needed in order to lower their free fatty acid content and peroxide value. It should be noted that the elevated peroxide levels may be due to the extended time lapse (approximately one month) which occurred between collection of the clay sample at the soybean oil processing plant and the SFE-processing of the clays at NCAUR.

Table 3

PROPERTER	or our	DITRACTED	FROM	SPENT	BLEACHING	CLAYS
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MAN	FFA PV (SI) (meg/Kgi)		Lovibend Color	Phospharus (ppm)	
ŧ	.46	1.8	44.0Y/2.7R	8	
2	.48	1.5	42.0Y/2.7R	.16	
2	.48	1.2	44.0Y/2.7R	.46	
4	.40	1.1	80.0Y/3.1R	.16	
7	.87	2.9	68.0Y/12.1R	.51	
•	.19	3.5	68.0Y/E.3R	.30	
10	.57	7.7	70.2Y/13.4R	2.00	

A visual examination of the SFE-extracted bleaching clays revealed that a reduction in color had been achieved when they were compared to the original product from the oil refinery. However, the SFE-treated clay still contained more pigment matter than the virgin clay. The bleaching power of treated and recycled clays has been studied by Ong and Sinkeldam (13), who found that the bleaching power was 50% less than that achieved by using fresh clay. For this reason, no further studies were attempted to assess the bleaching power of the SC-CO2 treated clays. Separation of the treated clays from the Hydromatrix dispersant could be easily performed by sieving methods, due to the large particle size difference between the spent clay and the Hydromatrix.

CONCLUSIONS

The above SFE results show that quantitative recovery of soybean oil from neutral and acidic spent bleaching clays is possible using CO2. Experiments conducted in both laboratory and

pilot plant extractors yielded variable oil recovery results, unless suitable precautions were taken to avoid fluid channeling and bed plugging. The use of a dispersing agent was found to enhance oil recovery and an improve extraction kinetics. An improved method was developed for determining residual oil levels by Soxhlet extraction. Finally, spent clays can be treated readily in a semi-continuous extraction mode, yielding an industrial-grade soybean oil and oil-free clay.

ACKNOWLEDGEMENTS

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DISCLAIMER

The mention of firm names or trade products does not imply that they are endorsed or recommended by the U.S. Dept. of Agriculture over other firms or similar products not mentioned.

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